Food and Drug Administration, HHS

§872.3590 Preformed plastic denture tooth.

- (a) *Identification*. A preformed plastic denture tooth is a prefabricated device, composed of materials such as methyl methacrylate, that is intended for use as a tooth in a denture.
 - (b) Classification. Class II.

§ 872.3600 Partially fabricated denture kit.

- (a) Identification. A partially fabricated denture kit is a device composed of connected preformed teeth that is intended for use in construction of a denture. A denture base is constructed using the patient's mouth as a mold, by partially polymerizing the resin denture base materials while the materials are in contact with the oral tissues. After the denture base is constructed, the connected preformed teeth are chemically bonded to the base.
- (b) Classification. Class II. The special controls for this device are FDA's:
- (1) "Use of International Standard ISO 10993 'Biological Evaluation of Medical Devices—Part I: Evaluation and Testing," and (2) "OTC Denture Reliners, Repair
- (2) "OTC Denture Reliners, Repair Kits, and Partially Fabricated Denture Kits."

 $[52~{\rm FR}~30097,~{\rm Aug.}~12,~1987,~{\rm as~amended}~{\rm at}~65~{\rm FR}~17144,~{\rm Mar.}~31,~2000]$

§872.3630 Endosseous dental implant abutment.

- (a) Identification. An endosseous dental implant abutment is a premanufactured prosthetic component directly connected to the endosseous dental implant and is intended for use as an aid in prosthetic rehabilitation.
- (b) Classification. Class II (special controls). The guidance document entitled "Class II Special Controls Guidance Document: Root-Form Endosseous Dental Implants and Endosseous Dental Implant Abutments" will serve as the special control. (See §872.1(e) for the availability of this guidance document.)

[69 FR 26304, May 12, 2004]

§ 872.3640 Endosseous dental implant.

(a) *Identification*. An endosseous dental implant is a device made of a mate-

rial such as titanium or titanium alloy, that is intended to be surgically placed in the bone of the upper or lower jaw arches to provide support for prosthetic devices, such as artificial teeth, in order to restore a patient's chewing function.

- (b) Classification. (1) Class II (special controls). The device is classified as class II if it is a root-form endosseous implant. The dental root-form endosseous dental implant is characterized by four geometrically distinct types: Basket, screw, solid cylinder, and hollow cylinder. The guidance document entitled "Class II Special Controls Guidance Document: Root-Form Endosseous Dental Implants Endosseous Dental Implant Abutments" will serve as the special control. (See §872.1(e) for the availability of this guidance document.)
- (2) Class III (premarket approval). The device is classified as class III if it is a blade-form endosseous dental implant.

 $[69\;\mathrm{FR}\;26304,\,\mathrm{May}\;12,\,2004]$

§872.3645 Subperiosteal implant material.

- (a) Identification. Subperiosteal implant material is a device composed of titanium or cobalt chrome molybdenum intended to construct custom prosthetic devices which are surgically implanted into the lower or upper jaw between the periosteum (connective tissue covering the bone) and supporting bony structures. The device is intended to provide support for prostheses, such as dentures.
 - (b) Classification. Class II.

$\S 872.3660$ Impression material.

- (a) Identification. Impression material is a device composed of materials such as alginate or polysulfide intended to be placed on a preformed impression tray and used to reproduce the structure of a patient's teeth and gums. The device is intended to provide models for study and for production of restorative prosthetic devices, such as gold inlays and dentures.
- (b) Classification. Class II (Special Controls).

[52 FR 30097, Aug. 12, 1987, as amended at 68 FR 19738, Apr. 22, 2003]